

## UNITED STATES DEPARTMENT OF COMMERCE Patent and Trademark Office

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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	AT	TORNEY DOCKET NO.
		¬ [	EX	KAMINER
		Г	ART UNIT	PAPER NUMBER
				:2
. IX **			DATE MAILED:	

Please find below and/or attached an Office communication concerning this application or proceeding.

**Commissioner of Patents and Trademarks** 

## Office Action Summary

Application No 09/384,959

Appli

(S)

Examiner

Richard Hutson

Group Art Unit

1652

Sasisekharan et al.



	ponsive to communication(s) filed on <u>Aug 2, 1999</u>			
This	action is FINAL.			
Since in ac	te this application is in condition for allowance except for formal mocordance with the practice under <i>Ex parte Quay</i> (835 C D 11, 45)	natters. <b>prosecution as to the merits is closed</b> 453 O G 213		
longer. f applicat	ened statutory period for response to this action is set to expire $\underline{}$ from the mailing date of this communication. Failure to respond value to become abandoned (35 U S C § 133). Extensions of time R 1.136(a).	within the period for response will cause the		
Disposi	ition of Claim			
ΧC	Claim(s) <u>1-57</u>	is/are pending in the applicat		
C	Of the above, claim(s)	is/are withdrawn from considerati		
C	Claim(s)	is/are allowed		
C	Claim(s)	is/are rejected		
Claim(s)				
	Claims <u>1-57</u>			
S T T T Priority	See the attached Notice of Draftsperson's Patent Drawing Review, The drawing(s) filed on	to by the Examiner.  is approveddisapproved  5 U S.C § 119(a)-(d).  rity documents have been  tional Bureau (PCT Rule 17 2(a))		
	ment(s) Notice of References Cited, PTO-892			
	nformation Disclosure Statement(s), PTO-1449, Paper No(s)			
Ir	nterview Summary, PTO-413			
	Notice of Draftsperson's Patent Drawing Review, PTO-948			
11				

--- SEE OFFICE ACTION ON THE FOLLOWING PAGES ---

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## DETAILED ACTION

## Election/Restriction

Restriction to one of the following inventions is required under 35 U.S.C. 121:

- Claims 1-21 and 52-54, drawn to a modified heparinase II protein, classified in class 435, subclass 200.
- Claim 22-29, drawn to a modified heparinase I protein, classified in class 435, subclass 200.
- III. Claims 30-34 and 46-49, drawn to a method of specifically cleaving a heparin-like glucosaminoglycan, classified in class 435, subclass 101.
- IV. Claims 35-44, drawn to a method of inhibiting angiogenesis, classified in class 424, subclass 94.61.
- V. Claims 45 and 514, drawn to a method of sequencing heparin and heparan sulfate. classified in class 435, subclass 101.
- VI. Claim 50, drawn to a method of inhibiting cellular proliferation, classified in class 424, subclass 94.61.
- VII. Claims 55-57, drawn to an isolated nucleic acid encoding a heparinase, classified in class 536, subclass 23.2.

The inventions are distinct, each from the other because of the following reasons:

Inventions I, II and VII are unrelated. Inventions are unrelated if it can be shown that they are

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not disclosed as capable of use together and they have different modes of operation, different functions, or different effects (MPEP § 806.04, MPEP § 808.01). In the instant case the modified heparinases of groups I and II and the nucleic acid encoding a heparinase of Group VII each comprise a chemically unrelated structure capable of separate manufacture, use and effect. The modified heparinases of groups I and II each comprise a different amino acid sequence and the DNA of Group VII is comprised of nucleic acid sequence. The DNA has other utility besides encoding protein such as a hybridization probe, and the proteins can be made synthetically. Additionally, the protein can be used to perform specific biological function(s) which are independent of the function(s) of the DNA molecule. The protein has other utility such as for the method of Group III.

Inventions I or II and III -VI are related as product and process of use. The inventions can be shown to be distinct if either or both of the following can be shown: (1) the process for using the product as claimed can be practiced with another materially different product or (2) the product as claimed can be used in a materially different process of using that product (M.P.E.P. § 806.05(h)). In the instant case the each of the different products of groups I or II can be used to for the different methods of group III-VI.

The nucleic acid of group VIII is unrelated to the method of groups III- VI as it is neither used nor made by the methods of groups III-VI.

The methods of groups III-VI are independent as they comprise different steps, utilize different products and produce different results.

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- Because these inventions are distinct for the reasons given above and the search required for Groups I-VII are not coextensive, restriction for examination purposes as indicated is proper. "For purposes of the initial requirement, a serious burden on the examiner may be prima facie shown if the examiner shows by appropriate explanation either separate classification, separate status in the art, or a different field of search as defined in MPEP 808.02." (see MPEP 803).
- 2. This application contains claims directed to the following patentably distinct species of the claimed invention: methods of using a heparinase II (claims 31, 46-51) or a heparinase I (claim 32) protein.

Applicant is required under 35 U.S.C. 121 to elect a single disclosed species for prosecution on the merits to which the claims shall be restricted if no generic claim is finally held to be allowable. Currently, claims 30, 33-45 are generic.

Applicant is advised that a reply to this requirement must include an identification of the species that is elected consonant with this requirement, and a listing of all claims readable thereon, including any claims subsequently added. An argument that a claim is allowable or that all claims are generic is considered nonresponsive unless accompanied by an election.

Upon the allowance of a generic claim, applicant will be entitled to consideration of claims to additional species which are written in dependent form or otherwise include all the limitations of an allowed generic claim as provided by 37 CFR 1.141. If claims are added after the election, applicant must indicate which are readable upon the elected species. MPEP \$809.02(a).

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Should applicant traverse on the ground that the species are not patentably distinct. applicant should submit evidence or identify such evidence now of record showing the species to be obvious variants or clearly admit on the record that this is the case. In either instance, if the examiner finds one of the inventions unpatentable over the prior art, the evidence or admission may be used in a rejection under 35 U.S.C. 103(a) of the other invention.

Applicant is advised that the reply to this requirement to be complete must include an election of the invention to be examined even though the requirement be traversed (37 CFR 1.143).

3. Applicant is reminded that upon the cancellation of claims to a non-elected invention, the inventorship must be amended in compliance with 37 CFR 1.48(b) if one or more of the currently named inventors is no longer an inventor of at least one claim remaining in the application. Any amendment of inventorship must be accompanied by a petition under 37 CFR 1.48(b) and by the fee required under 37 CFR 1.17(i).

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Richard Hutson whose telephone number is (703) 308-0066. The examiner can normally be reached on M-F from 7:30 to 4:00.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Ponnathapy Achutamurthy (Murthy), can be reached on (703) 308-3804. The fax number for Official Papers to Technology Center 1600 is (703) 305-3014.